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(21) International Application Number: PCT/EP98/03204 (22) International Filing Date: 29 May 1998 (29.05.98) (30) Priority Data: 97109619 30 May 1997 (30.05.97) RU (71) Applicant (for all designated States except US): ALFAR INTERNATIONAL LTD. [IE/IE]; 20 Clan William Terrace, Dublin 2 (IE). (72) Inventors; and (75) Inventors/Applicants (for US only): GORDEEV, Sergey Konstantinovich [RU/RU]; Apartment 27, pr. Rybatskij, 19/1, St.Petersburg, 193076 (RU). AVARBZ, Robert Gustavovitj [RU/RU]; Apartment 23, Korolenkostreet, 7, St.Petersburg, 191104 (RU). KRAVTJIK, Aleksandr Efimovitj [RU/RU]; Apartment 48, Statjek prospect, 11, St.Petersburg, 198097 (RU). KUKUSJKINA, Julia Aleksandrovna [RU/RU]; Apartment 143, Bolsjeoktinskij prospect, 14, St.Petersburg, 196160 (RU). SOKOLOV, Vasilij Vasilevitj [RU/RU]; Apartment 25, Tjernoje Kazatjestvostreet, 8, St.Petersburg, 198096 (RU). MAZA-EVA, Tatjana Vasilevna [RU/RU]; House 2, Apartment 135, Kosygin prospect, 30, St.Petersburg, 195208 (RU).		GRECHINSKAYA, Alla [RU/RU]; Apartment 22, Kuibushev Street, 20, St.Petersburg, 19746 (RU). (74) Agents: HYLTURNER, Jan-Olof et al.; Noréns Patentbyrå AB, P.O. Box 10198, S-100 55 Stockholm (SE). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: A METHOD FOR PRODUCING A POROUS CARBON ARTICLE AND AN ARTICLE PRODUCED THEREBY (57) Abstract <p>The present invention relates to a method for producing a porous carbon article comprising the steps of forming a workpiece with transport porosity and forming nanopores in said workpiece by thermochemical treatment. According to the invention the workpiece is formed as a rigid carbonaceous skeleton containing in its structure particles of one or more carbides, being selected and arranged in order to provide predetermined nanopore sizes, a predetermined volume of nanopores and a predetermined distribution of nanopores within the volume of the article.</p>		

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A METHOD FOR PRODUCING A POROUS CARBON ARTICLE AND AN ARTICLE PRODUCED THEREBY

TECHNICAL FIELD

5

The present invention relates to a method for producing a porous carbon article comprising the steps of forming a workpiece with transport porosity and forming nanopores in said workpiece by thermochemical treatment. The invention also
10 relates to an article produced by said method.

BACKGROUND TO THE INVENTION

From "Application of tubular articles in cryoadsorption
15 pumps//Carbon adsorbents and their application in industries.", Breslavets K.S et al, Moscow, Science publishers, 1983, p. 243, a method for producing a porous carbon article is known. The method comprises a step of forming or extrusion of a paste consisting of silicon carbide powder and commercial synthetic resins as a binder in order to produce a de-
20 sired article. In this case, transport porosity of the material is formed with pore size above 100 nm. Then a carbonization in an inert medium is carried out in order to strengthen the article and make its structure more uniform. Further the
25 article undergoes a thermochemical treatment by chlorine at 900 - 1000°C for transformation of carbide into carbon. In this step, in the volume of the article a nanoporous structure with pore size less than 10 nm is formed.

30 Use of a polymeric resin as a binder is an obstacle for obtaining high mechanical strength, because of the low mechanical strength of carbonized resin. Resin destruction accompanies formation of carbon which also takes part in the process of forming nanoporosity, but the size of this porosity is
35 practically uncontrollable. As a result, it is impossible to produce materials with predetermined adsorption properties with the known method.

An article produced by means of the known method is a carbon material binded with products of resin carbonization with porosity of 65 to 75 vol%. In this case, a part of the pores, 30-32 vol% are transport pores having size greater than 100 nm, while other pores have size less than 10 nm.

Application of articles obtained by the known method is restricted because it gives no possibility to obtain controllable size of pores as well as controllable volumetric content of both transport porosity and nanoporosity.

A number of so called activated carbons with a high content of nanoporosity is known, but the pore size distribution for these materials is very wide and uncontrolled, c.f. "Carbon", John Wiley & Son, N.Y. 1988, USA.

It is thus a need for a method in which the porosities of a porous carbon material which comprise two types of pores can be controlled. The two types are pores of a size less than 10 nm providing adsorption ability and pores of a size greater than 100 nm providing transportation of a component to the pores taking active part in the adsorption process. Articles produced by such a method can be used in different fields of technology connected with adsorption and absorption processes, such as selective absorption of a component from a liquid or gas, electrochemical electrodes, in medicine technologies, etc.

The object of the present invention is to make it possible to produce carbon porous articles with predetermined transport porosity and predetermined nanopore sizes, volume and distribution throughout the volume of the article.

SUMMARY OF THE INVENTION

This object is achieved by a method for producing a porous carbon article comprising the steps of forming a workpiece with transport porosity and forming nanopores in said workpiece by thermochemical treatment, which is characterized in that the workpiece is formed as a rigid carbonaceous skeleton containing in its structure particles of one or more carbides, being selected and arranged in order to provide predetermined nanopore sizes, a predetermined volume of nanopores and a predetermined distribution of nanopores within the volume of the article. By this method materials having controlled and predetermined nanopores, an optimal ratio between volumes of transport pores and nanopores, high mechanical strength and complicated shapes can be produced.

In a preferred embodiment carbides from III, IV, V or VI group of Mendeleev's Periodic System are selected as carbon precursor. For carbides forming carbons having an assumed slot-like structure, the carbides are selected on the basis of dependence of specified nanopore size on physical and chemical constants of the carbides;

$$X = Z \cdot (1-R) / R$$

where X = specified size of nanopores, nm;

$$Z = 0.65-0.75 \text{ nm};$$

$$R = v M_C \rho_K / M_K \rho_C$$

where M_C - molecular mass of carbon, g/mole;

M_K - molecular mass of carbide, g/mole;

ρ_K - density of carbide, g/ccm;

ρ_C - density of carbon, g/ccm;

v - number of carbon atoms in carbide molecule.

The formulation of carbide particle mixture is chosen in dependence of desired distribution of nanopores by sizes using the relationship;

$$\Psi_i = K_i \varphi_i / \sum_{i=1}^n K_i \varphi_i$$

where Ψ_i - volumetric part of nanopores with size X_i in total volume of nanopores;

φ_i - volumetric part of i -th carbide in particle mixture;

n - number of carbides;

$$K_i = 1 - v M_C \rho_{ki} / M_{ki} \rho_C$$

where M_C - molecular mass of carbon, g/mole;

M_{ki} - molecular mass of i -th carbide, g/mole;

ρ_{ki} - density of i -th carbide, g/ccm;

ρ_C - density of carbon, g/ccm;

v - number of carbon atoms in carbide molecule.

The forming step comprises forming an intermediate body of particles of one or more carbides and subsequent treating of this body in a medium of gaseous hydrocarbon or hydrocarbon mixtures at a temperature exceeding the decomposition temperature for the hydrocarbon or hydrocarbons until the mass of the intermediate body has increased at least 3% and the intermediate body is formed with a porosity of 30-70 vol%, preferably 35-50 vol%. Furthermore, the intermediate body is formed with a porosity determined by the following relationship;

$$\varepsilon_0 = [1 - v_{np} / \sum_{i=1}^n K_i \varphi_i] \cdot 100$$

where ε_0 - porosity of intermediate body, vol%;

φ_i - volumetric part of i-th carbide in powder mixture;
 V_{np} - predetermined volumetric part of nanopores in
 final article.

$$K_i = 1 - vM_c\rho_{ki}/M_{ki}\rho_c$$

where M_c - molecular mass of carbon, g/mole;
 M_{ki} - molecular mass of i-th carbide, g/mole;
 ρ_{ki} - density of i-th carbide, g/ccm;
 ρ_c - density of carbon, g/ccm;
 v - number of carbon atoms in carbide molecule,

and the treatment in a medium of hydrocarbon or hydrocarbons
 is carried out until the mass of the intermediate body has
 changed according the following relationship;

$$\Delta m = Q(\varepsilon_0 - v_{tr})/(1 - \varepsilon_0)$$

where Δm - relative change of intermediate body mass, g/g;
 ε_0 - porosity of intermediate body, vol%;
 V_{tr} - predetermined volumetric content of transport
 pores, vol%;

$$Q = \rho_c/\rho_{mix}$$

where ρ_c - density of carbon, g/ccm:
 ρ_{mix} - density of carbides mixture, g/ccm.

The intermediate body can be formed by pressing. Other well
 known forming methods, such as slip casting, tape casting,
 slurry casting and injection moulding can of course also be
 used. Natural gas is used as a mixture of hydrocarbons and
 the treating in hydrocarbon medium is carried out at 750-
 950°C.

Alternatively is at least one of the hydrocarbons used during the treatment of the intermediate body in hydrocarbons medium selected from the group of acetylene, methane, ethane, propane, pentane, hexane, benzene and their derivatives and the
5 treating in hydrocarbon medium is carried out at 550-1200⁰C.

The thermochemical treatment is carried out in a medium of gaseous halogens creating volatile halogenides, such as chlorides, at 350-1200⁰C.

10

The particles of carbide or carbides of which the intermediate body is formed are arranged uniformly or nonuniformly throughout its volume.

15

The present invention relates also to a porous carbon article having nanopores and transport pores, which is characterized in that the volume of transport pores is 10-55% and the volume of nanopores is 15-50%. The nanopores have sizes in the range of 0.6-3.5 nanometers and are distributed uniformly or
20 nonuniformly throughout the volume of the article.

The following terms are currently used:

25

- for porosity with pore size of greater than 100 nm - transport porosity, or macroporosity;
- for porosity with pore size of less than 10 nm - nanoporosity.

These terms are used here to reveal the essence of the present invention.

30

BRIEF DESCRIPTION OF THE DRAWING

The present invention will now be described with reference to the following figures, of which;

35

Fig. 1 shows a table of the properties of materials produced in examples 1-3, and

Figs. 2-4 disclose porosimetry data for the samples of examples 1-3.

5 DESCRIPTION OF PREFERRED EMBODIMENTS OF THE INVENTION

The method according to the invention comprises the following steps:

- 10 1) Forming a workpiece with transport porosity using particles of a carbide or carbides of elements from III, IV, V and VI groups of Mendeleev's Periodic System, in the form of a rigid carbonaceous skeleton containing in its structure particles of a carbide or carbides selected from the said groups
15 and arranged in a predetermined order providing formation in the subsequent steps desired transport porosity and nanoporosity by sizes, volume and distribution of pores throughout the volume of the article;
- 20 2) Formation of nanoporosity throughout the volume of a workpiece obtained in the 1st step by thermochemical treatment of the said workpiece in gaseous halogens, such as chlorine, at elevated temperatures in the range of 350 to 1200°C, preferably 500-1100°C.

25 Current notions of carbon materials structure point out that nanopores generated during the thermochemical treatment process are formed by ordered or disordered graphite planes of carbon, which for simplicity might be considered as shaped as
30 slots, the width of the latter depending on type of carbide used for forming of the workpiece with transport porosity.

These theoretical ideas are in good agreement with experimental data which allowed the inventors to disclose the following
35 dependence for carbon materials having such structure:

$$X = Z \cdot (1-R)/R \quad (1)$$

where X-predetermined size of nanopores, nm;

- 5 Z - experimental factor established for a number of carbide structures of elements from III, IV, V and VI groups of Mendeleev's Periodic System as 0.65-0.75 nm;

$$R = vM_c\rho_k/M_k\rho_c$$

10

where M_c - molecular mass of carbon, g/mole;

M_k - molecular mass of carbide, g/mole;

ρ_k - density of carbide, g/ccm;

ρ_c - density of carbon, g/ccm;

15

v - number of carbon atoms in carbide molecule.

A series of preliminary experiments made it possible to choose a necessary carbide to obtain in practice a predetermined size of nanopores.

20

Particles of a chosen carbide (powder) are formed into an intermediate body with porosity in the range of 30-70 vol% by any known method, e.g. by pressing with or without temporary binder, slip casting, slurry casting. Final step of forming, which results in production of a workpiece with a high mechanical strength and a desired transport porosity, is a treating of the intermediate body in a medium of gaseous hydrocarbon or hydrocarbons mixture at a temperature above their decomposition temperature.

30

It is possible to use natural gas and/or at least a hydrocarbon selected from the group comprising acetylene, methane, ethane, propane, pentane, hexane, benzene and their derivatives.

35

Under these conditions a decomposition of hydrocarbon occurs by reaction;



with deposition of the generated pyrocarbon on the surface and in the pores of intermediate body volume.

The specified range of initial porosity is based on the fact that at a porosity below 30% it is difficult to obtain sufficient volume of transport pores in the article providing access of adsorptive to nanopores where adsorption process occurs and at a porosity above 70% the article does not possess satisfactory mechanical strength.

The value of 35-50 vol% is preferable because it is easily achieved by any available method of workpiece forming and it assures an optimal relation between volumes of transport pores and nanopores in the article.

The size and distribution of the transport pores can be controlled by selecting appropriate particle sizes and particle distribution. The amount of possible particle packing due to the forming process will of course also influence the porosity of the work piece.

Calculation of concrete value of intermediate body porosity necessary to obtain a predetermined volume of nanopores, is carried out using the following expression:

$$\varepsilon_0 = [1 - v_{np} / \sum_{i=1}^n K_i \varphi_i] \cdot 100 \quad (3)$$

where ε_0 - porosity of intermediate body, vol%;

φ_i - volumetric part of i-th carbide in powder mixture;

V_{np} - predetermined volumetric part of nanopores in final article.

$$K_i = 1 - vM_c\rho_{ki}/M_{ki}\rho_c$$

5

where M_c - molecular mass of carbon, g/mole;

M_{ki} - molecular mass of i-th carbide, g/mole;

ρ_c - density of carbon, g/ccm;

ρ_{ki} - density of i-th carbide, g/ccm;

10

v - number of carbon atoms in carbide molecule;

n - number of carbides in the mixture.

15

Duration of treating in the said medium is controlled by measuring the mass of the article. When the mass has changed by at least 3%, the strength is already sufficient for use of the article as adsorption element, capacitor electrode or chromatography membrane, for instance.

20

The process is usually completed when the mass is changed by 3-20%, thus providing necessary strength of the article and its transport porosity. Lower and upper limits are determined by use of carbides from said groups with different densities.

25

In practice an experimental expression is used allowing for a given type of carbide at predetermined strength properties to obtain necessary value of transport porosity which, depending on active agent in the pores, can determine kinetics of the process. This expression is as follows:

30

$$\Delta m = Q (\varepsilon_0 - V_{tr}) / (1 - \varepsilon_0); \quad (4)$$

where Δm - relative change of intermediate body mass, g/g;

ε_0 - porosity of intermediate body, vol%;

35

V_{tr} - predetermined volumetric content of transport pores, vol%;

$$Q = \rho_c / \rho_{\text{mix}}$$

where ρ_c - density of carbon, g/ccm:

ρ_{mix} - density of carbides mixture, g/ccm.

5

In order to obtain an article with nanopores of different sizes, making it possible to realize selective filtration and adsorption, more than one carbide should be chosen. For this goal the formula (1) or an experimentally determined pore size value is used and the following dependence, confirmed experimentally, allows an determination of the part of each carbide in the mixture necessary to manufacture such an article;

10

15

$$\Psi_i = K_i \varphi_i / \sum_{i=1}^n K_i \varphi_i \quad (5)$$

where ψ_i - volumetric part of nanopores with size X_i in total volume of nanopores;

φ_i - volumetric part of i-th carbide in powder mixture;

20

n - number of carbides;

$$K_i = 1 - v M_c \rho_{ki} / M_{ki} \rho_c$$

where M_c - molecular mass of carbon, g/mole;

25

M_{ki} - molecular mass of i-th carbide, g/mole;

ρ_{ki} - density of it-h carbide, g/ccm;

ρ_c - density of carbon, g/ccm;

v - number of carbon atoms in carbide molecule.

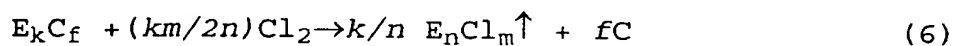
30

In order to obtain a uniform distribution of nanopores throughout the article volume a mixture is formed with uniformly distributed powders of various carbides in it (homogeneous mixture); if it is necessary to obtain nanopores distributed throughout the volume in a desired order a mix-

ture is prepared by means of any known method with particles distributed according to the desired order, e.g. layerwise.

5 After completed forming, a workpiece is obtained as a rigid carbonaceous skeleton with transport porosity formed in its volume allowing to obtain in the step of thermochemical treatment uniform nanopores of a predetermined size.

10 In order to form nanoporosity the obtained workpiece is subjected to thermochemical treatment by chlorine at 500-1100°C. Nanoporosity is formed at removal of volatile chlorides of carbide-forming elements in accordance with reaction:



15

where $E_k C_f$ - primary carbide;

k, f, n, m - stoichiometric coefficients.

20 The treatment is carried out until mass change of the workpiece has stopped.

A finished article produced by the described method has a predetermined shape and size, and its structure is a porous carbon skeleton with transport porosity of 10-55% obtained in
25 the step of forming and nanoporosity of 15-50% volume. The article comprises one or several types of nanopores and each type is being characterized with narrow distribution by size. Carbon content in the skeleton is more than 95 wt%, preferably 99 wt%, i.e., practically, the obtained article consists
30 of pure carbon and has considerable strength allowing to increase its life time and expand application range under conditions when shape maintaining during operation is necessary.

35 As a result of selecting appropriate carbides and accomplishment of forming under conditions determined beforehand by means of relationships established by the inventors, a fin-

ished article is obtained with nanopore sizes, volume and distribution corresponding to those of the object of the article operation.

- 5 Among possible forming methods to realize the said method pressing, slip casting, tape casting and slurry casting can be named.

10 A formed intermediate body is treated in a medium of at least one hydrocarbon selected from the group comprising acetylene, methane, ethane, propane, pentane, hexane, benzene and their derivatives. When using hydrocarbons from the said group an optimal temperature range is 550-1200°C, the decomposition
15 temperatures for these hydrocarbons falling within this range. It is also possible to use natural gas and in this case it is expedient to keep temperature in the range of 750-950°C.

A halogenation is carried out just like in the known method,
20 with the temperature being selected in the range of 350-1200°C, depending on the nature of initial carbides and the formed volatile halogenides. Under these conditions volatile halogenides of carbide-forming elements are completely re-
25 moved out of the article according to a reaction similar to reaction (6). However, only halogens and halogenides which do not react with carbon under the prevailing temperature conditions may be used.

The claimed concept is further elucidated with the following
30 examples.

Example 1.

An example of producing an article in the tablet form with
35 sizes $d = 20$ mm, $h = 5$ mm, with nanopore size 0.8 nm and nanopore volume 0.3 ccm/ccm uniformly distributed throughout the article volume, suitable for forming on its surface a

double electric layer of high capacitance in electrolyte solutions.

To produce an article on the basis of beforehand obtained dependence (1) for $X = 0.8$ nm titanium carbide powder was chosen. By substitution of the values of molecular mass and density of titanium carbide and carbon ($M_C = 12$ g/mole; $\rho_C = 2.2$ g/ccm; $\rho_K = \rho_{TiC} = 4.92$ g/ccm; $M_K = M_{TiC} = 59.88$ g/mole) in the formula (1) the following is obtained:

$$R = 12 \cdot 4.92 / 59.88 \cdot 2.2 = 0.448, \quad X = Z(1 - 0.448) / 0.448 = 1.232Z \text{ nm};$$

Thus when Z is in the range of 0.65-0.75, the nanopore size of the produced carbon materials will be in the range of 0.8-0.92.

In order to obtain the predetermined volume of nanopores ($V_{np} = 0.3$ ccm/ccm), prior to pressing a needed porosity of the intermediate body is determined by the relationship (3):

where $\varphi_i = 1$, $n = 1$, as follows

$$\varepsilon_0 = [1 - 0.3 / (1 - 0.448)] \cdot 100 = 46\%$$

Amount of TiC powder necessary to produce an intermediate body which has the predetermined sizes and the obtained value of porosity is calculated by the following dependence:

$$m = \rho_K (100 - \varepsilon_0) \cdot V / 100$$

where V - article volume, $V = (\pi d^2 / 4) \cdot h$, ccm;

d - workpiece diameter, 2 cm;

h - workpiece height, 0.5 cm;

hence:

$$m = 4.92 (100 - 46) (3.14 \cdot 2^2 / 4) \cdot 0.5 = 5.01g$$

The needed mass change of the workpiece during pyrocarbon deposition is calculated by formula (4), assuming a transport porosity of 35 vol%

Then, $\Delta m = [0.4476(46-35)/(100 - 46)] \cdot 100 = 9.1\%$

5

A mixture is prepared using 5.01 g of TiC powder with a size of the particles of 20 μm . Ethyl alcohol is added in the amount of 10% of the mass of the mixture. Then, an intermediate body is formed by pressing on a hydrostatic press machine (P-125) at 30 ± 1 Mpa pressure. After the pressing, the intermediate body is dried at $150 \pm 10^\circ\text{C}$ during 1-1.5 hour until complete removal of temporary binder.

15 This is followed by pyrocarbon deposition on the workpiece by means of heat treatment in natural gas medium at atmospheric pressure in a quartz continuous reactor at 850°C during 12 hours until change of mass by 9.1%.

20 Then, the sample is chlorinated. The chlorination is carried out in a isothermal quartz reactor at 650°C during 4 hours. Then a blow-through of the reactor with argon at a temperature of 800°C is carried out to remove excessive chlorine out of the reactor zone and the internal surface of the sample.

25

Properties of the obtained material are presented in Table 1. From this table it is evident that the measured peak value of the nanopore size measured by gas porosimetry correspond to the calculated value.

30

Two articles produced according to Example 1 were saturated with 20% KOH solution by boiling and placing them in an electrolyte solution (20% KOH). Opposite by sign potential was applied to each of the articles to form a double electric layer in the material nanopore volume. In this case the spe-

35

cific electrical capacitance of the double electric layer formed in the material was 37.8 F/g.

Example 2.

5

The example of producing an article in a tablet form of $d=30$ mm, $h=3$ mm with a nanopore volume 0.4 ccm/ccm uniformly distributed throughout the article volume suitable for adsorption of organic substances out of a powder of Mo_2C was chosen.

10

In order to produce the predetermined nanopore volume ($v_{np}=0.4$ ccm/ccm) before pressing the needed porosity of the intermediate body was calculated using expression (3), where

15 $M_{\text{Mo}_2\text{C}}=203,88$ g/mole; $\rho_{\text{Mo}_2\text{C}}=8.91$ g/ccm, $R=0.238$; $\phi_1=1$, $n=1$, as follows: $\varepsilon_0=48\%$

The amount of molybden carbide powder necessary to produce the intermediate body having the predetermined porosity was

20 calculated as in Example 1. In this case the powder article weight was calculated to: $m=9.82$ g

The needed mass change of the workpiece during pyrocarbon deposition was calculated by formula (4), assuming a transport porosity of 40vol%, to $\Delta m=3.8\%$.

25

Mixture preparation, pyrocarbon deposition and chlorination were carried out as in Example 1.

30 Properties of the obtained material are presented in Table 1. From this table it is evident that the calculated nanopore size of 1.95-2.25 nm differs from the measured peak value according to figures 3 and 4 which was 3.5 nm. This indicates that Mo_2C probably does not follow the model structure for

35 dependence (1).

The produced article was placed in an exsiccator containing isooctane vapours and was kept there during 24 hours. Then, the article was weighed to determine the amount of adsorbed isooctane, which was 0.95 ccm/g.

5

Example 3.

An example of producing an article of diameter 30 mm and height 5 mm with nanopores of calculated size 0.8 nm and 2.1 nm uniformly distributed throughout the article volume. In order to obtain the article on the basis of beforehand obtained dependence (1) for $X_1 = 0.8$ nm titanium carbide powder was chosen, for $X_2 = 3.5$ nm - molybdenum carbide powder in accordance with the measured value in Example 2.

15

In order to provide equal volumetric content of both sizes nanopores a mixture is used containing 40 vol% molybdenum carbide and 60 vol% titanium carbide, that is determined by formula (5). The needed amount of these carbides is calculated as follows:

20

$$\rho_{\text{mix}} = \varphi_1 \cdot \rho_1 + \varphi_2 \cdot \rho_2$$

where φ_1, φ_2 - volumetric content of titanium carbide and molybdenum carbide, correspondingly $\varphi_1 = 0.4$, $\varphi_2 = 0.6$);

25

ρ_1, ρ_2 - density of titanium carbide and molybdenum carbide, correspondingly ($\rho_1 = 8.91$ g/ccm, $\rho_2 = 4.92$ g/ccm);

hence $\rho_{\text{mix}} = 0.4 \cdot 8.91 + 0.6 \cdot 4.92 = 6.52$ g/ccm,

30

hence mass part of molybdenum carbide:

$$\alpha_1 = 0.4 \cdot 8.91 / 6.52 = 0.55 \text{ wt.fraction};$$

35

of titanium carbide

$$\alpha_2 = 0.6 \cdot 4.92 / 6.52 = 0.45 \text{ wt.fraction}$$

The mixture is prepared and pressed under conditions of Example 1.

In order to obtain an article of the specified shape and sizes a weight should be calculated according to the following dependence:

$$m = \rho_{\text{mix}}(100 - \varepsilon_0) \cdot V / 100$$

where ρ_{mix} - density of carbide mixture;

ε_0 - porosity of intermediate body; %vol;

V - article volume, ccm;

d - article diameter, 3 cm;

h - article height, 0.5 cm.

The needed porosity of an intermediate body is chosen according to relation (3).

By substitution of the said values at specified total volume of nanopores 0.4 ccm/ccm we obtain

$$\varepsilon_0 = [100 - 0.4 / ((1 - 12 \cdot 8.91 / 2.2 \cdot 203.88) 0.4 + (1 - 12 \cdot 4.92 / 2.2 \cdot 59.88) 0.6)] 100 \approx 37 \text{ vol\%}$$

hence necessary mass of the weight:

$$m = 6.52(100 - 37) \cdot (3.14 \cdot 5^2 / 4) \cdot 0.2 / 100 = 16.1 \text{ g}$$

The obtained intermediate body is heat treated under conditions of Example 1. Introduction of pyrocarbon is carried out under conditions of Example 1 until change of article mass by 7% which is determined by formula (4) under condition:

$$V_{\text{tr}} = 20 \text{ vol\%}$$

$$\Delta m = 0.337(37-20)/(100 - 37) \cdot 100 = 9.1\%$$

Chlorination of the obtained workpiece is carried out under conditions of Example 1.

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Properties of samples produced in Examples 1 - 3 are presented in Table 1. From Table 1 it seems as Mo_2C results in a different carbon nanopore structure than TiC .

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The produced article was placed in an exsiccator containing vapours of carbontetrachloride and was kept there during 24 hours. Then the article was weighed to determine the amount of absorbed carbontetrachloride which was 0.61 ccm/g.

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Notes:

1) Total volume of pores is determined by hydrostatic method according to GOST 473.4-81.

2) Nanopore volume is determined by exsiccator method by adsorption of benzene under static conditions, see "Fundamentals of adsorption technology." Keltsev N.V., Moscow, Chemistry publishers, 1984, p. 33.

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3) Transport pore volume is determined by formula

$$V_{tr} = V_{\Sigma} - V_{np}.$$

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4) Size of nanopores is determined by means of mercury and gas porosimetry (Micromeritics Auto Pore III and Micromeritics ASAP 2010, respectively). Data are shown in Figures 2-4. Legend Hg denotes mercury porosimetry intrusion data, legend BJH denotes gas porosimetry desorption data analysed by the BJH method, and legend Micro denotes gas porosimetry data analysed by the Horvath-Kawazoe method.

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The presented data allows one to draw the conclusion that a new method for producing a porous carbon article comprising transport pores and nanopores with controllable sizes and distribution of nanopores throughout its volume as well as

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volumetric content of both types of porosity has been developed. The articles according to the invention can find wide application for adsorption and microdosage of substances, purification and separation of cryogenic liquids and gas mixtures, as high-porous electrode materials etc. owing to presence of porosity of desired sizes.

By the inventive method it is possible to produce nanopore volume and size or sizes by a mechanism independent from the mechanism for producing transport porosity in the materials produced, thereby making it possible to control purposely parameters of their porous structure. At development of adsorption materials, for instance, the following parameters can be optimized when using the present invention:

- 1) adaptability to manufacture of devices working components made of these materials;
- 2) optimal relationship between volumes of transport pores and nanopores which provide effective adsorption;
- 3) mechanical strength;
- 4) increased heat conductivity allowing to use these materials in cryo-adsorption evacuation elements.

Furthermore, the present method, besides the advantages pointed out, allows production of articles of complex shapes, in particular, of shapes impossible to obtain by any other known method, with minimum machining required. Owing to high mechanical strength the articles according to the invention can be used under conditions demanding maintenance of their shape.

CLAIMS

1. A method for producing a porous carbon article comprising the steps of forming a workpiece with transport porosity and forming nanopores in said workpiece by thermochemical treatment, characterized in that the workpiece is formed as a rigid carbonaceous skeleton containing in its structure particles of one or more carbides, being selected and arranged in order to provide predetermined nanopore sizes, a predetermined volume of nanopores and a predetermined distribution of nanopores within the volume of the article.

2. A method according to Claim 1, characterized in that carbides from III, IV, V or VI group of Mendeleev's Periodic System are selected as carbon precursor.

3. A method according to Claim 1 or 2, characterized in that for carbides forming carbons having a slot-like structure, the carbides are selected on the basis of dependence of specified nanopore size on physical and chemical constants of the carbides;

$$X = Z * (1-R) / R \quad (1)$$

25

where X = specified size of nanopores, nm;

$Z = 0.65-0.75$ nm;

$$R = vM_C\rho_K / M_K\rho_C$$

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where M_C - molecular mass of carbon, g/mole;

M_K - molecular mass of carbide, g/mole;

P_K - density of carbide, g/ccm;

P_C - density of carbon, g/ccm;

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v - number of carbon atoms in carbide molecule.

4. A method according to any one of Claims 1-3, characterized in that the formulation of carbide particle mixture is chosen in dependence of desired distribution of nanopores by sizes using the relationship;

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$$\Psi_i = K_i \varphi_i / \sum_{i=1}^n K_i \varphi_i$$

where Ψ_i - volumetric part of nanopores with size X_i in total volume of nanopores;

φ_i - volumetric part of i-th carbide in particle mixture;

10

n - number of carbides;

$$K_i = 1 - v M_C \rho_{ki} / M_{ki} \rho_C$$

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where M_C - molecular mass of carbon, g/mole;

M_{ki} - molecular mass of i-th carbide, g/mole;

ρ_{ki} - density of i-th carbide, g/ccm;

ρ_C - density of carbon, g/ccm;

v - number of carbon atoms in carbide molecule

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5. A method according to any one of Claims 1-4, characterized in that the forming step comprises forming an intermediate body of particles of one or more carbides and subsequent treating of this body in a medium of gaseous hydrocarbon or hydrocarbon mixtures at a temperature exceeding the decomposition temperature for the hydrocarbon or hydrocarbons until the mass of the intermediate body has increased at least 3%.

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6. A method according to any one of claims 1-5, characterized in that the intermediate body is formed with a porosity of 30-70 vol%, preferably 35-50 vol%.

7. A method according to any one of Claims 1-6, characterized in that the intermediate body is formed with a porosity determined by the following relationship;

5

$$\varepsilon_0 = [1 - v_{np} / \sum_{i=1}^n K_i \varphi_i] * 100$$

where ε_0 - porosity of intermediate body, vol%;

φ_i - volumetric part of i-th carbide in powder mixture;

10 v_{np} - predetermined volumetric part of nanopores in final article.

$$K_i = 1 - v_{M_C} \rho_{ki} / M_{ki} \rho_C$$

15 where M_C - molecular mass of carbon, g/mole;

M_{ki} - molecular mass of i-th carbide, g/mole;

ρ_{ki} - density of i-th carbide, g/ccm;

ρ_C - density of carbon, g/ccm;

v - number of carbon atoms in carbide molecule

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8. A method according to any one of Claims 1-7, characterized in that the treatment in a medium of hydrocarbon or hydrocarbons is carried out until the mass of the intermediate body has changed according the following relationship;

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$$\Delta m = Q(\varepsilon_0 - v_{tr}) / (1 - \varepsilon_0)$$

where Δm - relative change of intermediate body mass, g/g;

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ε_0 - porosity of intermediate body, vol%;

v_{tr} - predetermined volumetric content of transport pores, vol%;

$$Q = \rho_C / \rho_{mix}$$

where ρ_c - density of carbon, g/ccm:

ρ_{mix} - density of carbides mixture, g/ccm.

- 5 9. A method according to any one of Claims 1-8,
c h a r a c t e r i z e d i n that the intermediate body is
formed by pressing.
- 10 10. A method according to any one of Claims 1-8,
c h a r a c t e r i z e d i n that the intermediate body is
formed by slip casting, tape casting or slurry casting.
- 15 11. A method according to any one of Claims 1-10,
c h a r a c t e r i z e d i n that natural gas is used as a
mixture of hydrocarbons.
- 20 12. A method according to any one of Claims 1-11,
c h a r a c t e r i z e d i n that the treating in hydrocar-
bon medium is carried out at 750-950⁰C.
- 25 13. A method according to any one of Claims 1-10,
c h a r a c t e r i z e d i n that at least one of the hy-
drocarbons used during the treatment of the intermediate body
in hydrocarbons medium is selected from the group of acety-
lene, methane, ethane, propane, pentane, hexane, benzene and
their derivatives.
- 30 14. A method according to any one of Claims 1-10 or 13,
c h a r a c t e r i z e d i n that the treating in hydrocar-
bon medium is carried out at 550-1200⁰C.
- 35 15. A method according to any one of Claims 1-14,
c h a r a c t e r i z e d i n that the thermochemical treat-
ment is carried out in a medium of gaseous halogens, such as
chlorine.

16. A method according to any one of Claims 1-15,
c h a r a c t e r i z e d i n that the thermochemical treat-
ment is carried out at 350-1200⁰C.
- 5 17. A method according to any one of Claims 1-16,
c h a r a c t e r i z e d i n that the the particles of
carbide or carbides of which the intermediate body is formed
are arranged uniformly throughout its volume.
- 10 18. A method according to any one of Claims 1-16,
c h a r a c t e r i z e d i n that that the the particles of
carbide or carbides of which the intermediate body is formed
are arranged nonuniformly throughout its volume.
- 15 19. A porous carbon article having nanopores and transport
pores, c h a r a c t e r i z e d i n that the volume of
transport pores is 10-55%.
- 20 20. An article according to Claim 19,
c h a r a c t e r i z e d i n that the volume of nanopores
is 15-50%.
- 25 21. An article according to Claim 19 or 20,
c h a r a c t e r i z e d i n that the nanopores have peak
value sizes of 0.8 and/or 3.5 nanometer.
- 30 22. An article according to any one of Claims 19-21,
c h a r a c t e r i z e d i n that the nanopores are dis-
tributed uniformly throughout the volume of the article.
23. An article according to any one of Claims 19-21,
c h a r a c t e r i z e d i n that the nanopores are dis-
tributed nonuniformly throughout the volume of the article.

24. An article according to Claim 20,
c h a r a c t e r i z e d i n having a specific electrical
capacitance of at least 30 F/g when used as electrode mate-
5 rial in a double electric layer capacitor.

1 / 2

Table 1

Ex. No	Carbide mixture composition (wt%)	Total volume of pores ¹⁾ V_{tot} (cc/cc)	Nanopore volume, ²⁾ V_{np} (cc/cc)	Transport pore volume ³⁾ , V_{tr} (cc/cc)	Size of pores, calculated ⁴⁾ (nm)		Size of pores, measured ⁵⁾ (nm)	
					X_1	X_2	X_1	X_2
1	100% TiC	0,67	0,32	0,35	0.74- 0.85		0.8	
2	100% Mo2C	0,80	0,39	0,41		1.95- 2.25		3.5
3	45% TiC + 55% Mo2C	0.62	0.41	0.21	0.74- 0.85	1.95- 2.25	0.8	3.5

FIG. 1

Carbon material from TiC

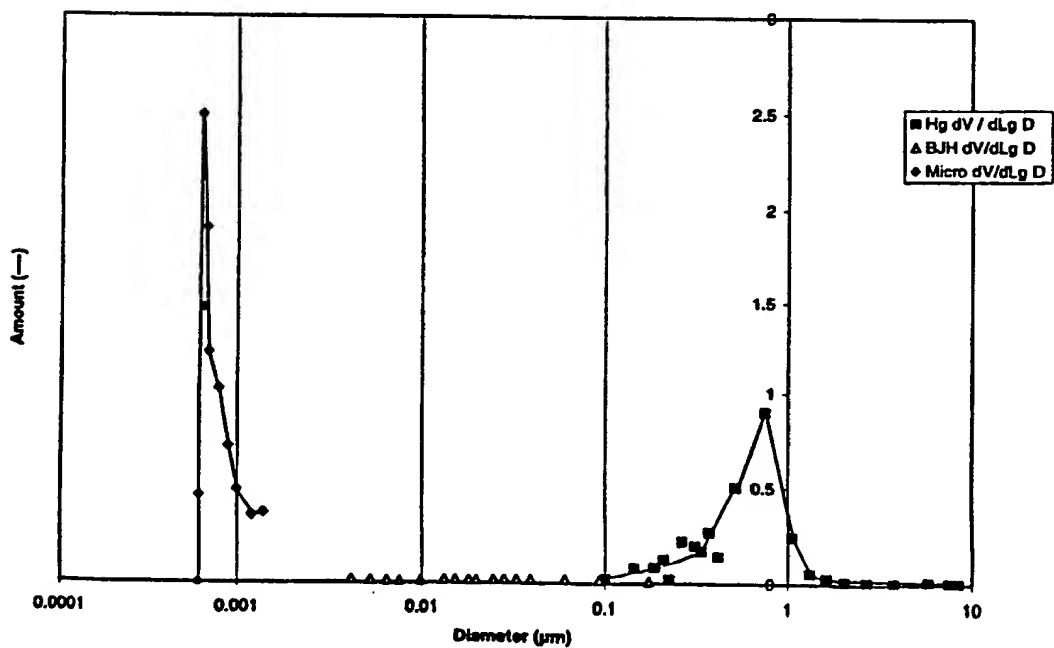


FIG. 2

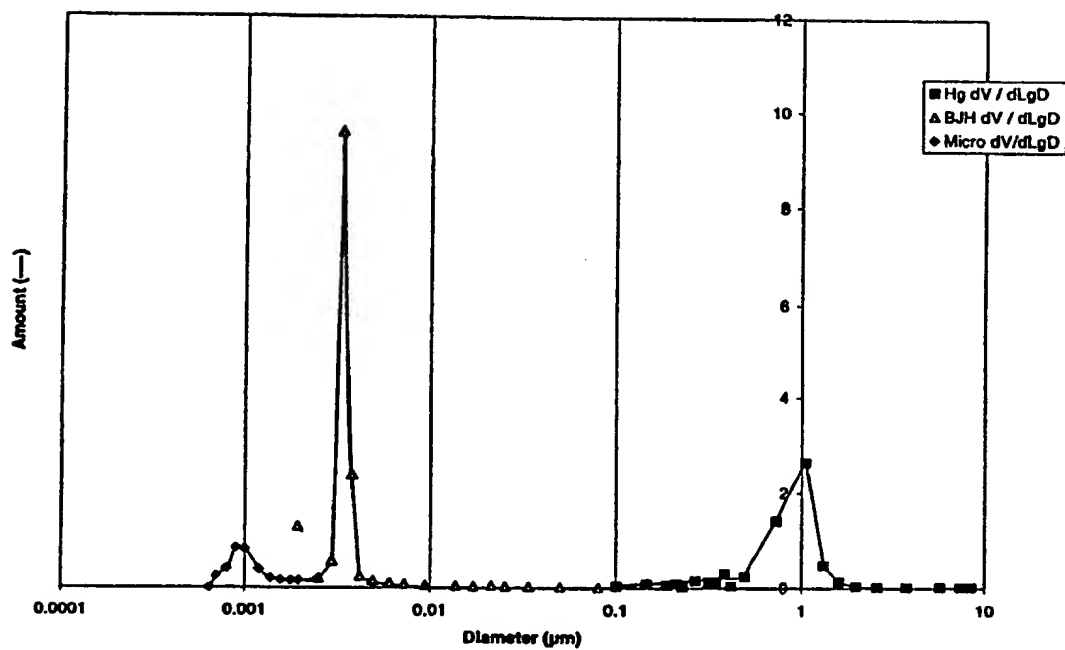
Carbon material from Mo₂C

FIG. 3

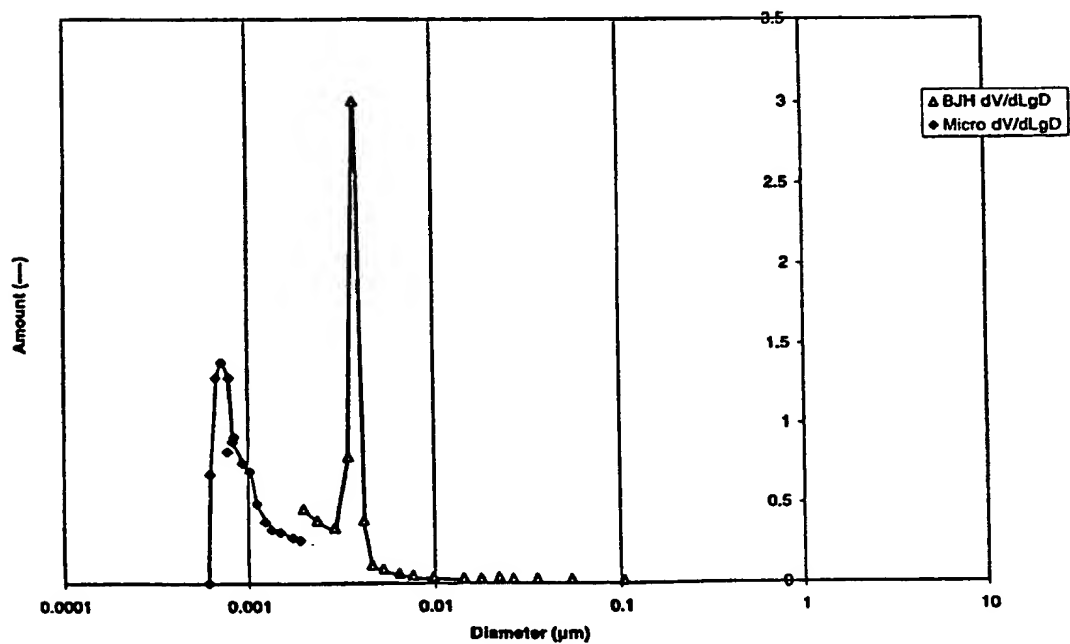
Carbon from TIC and Mo₂C

FIG. 4

INTERNATIONAL SEARCH REPORT

International Application No.
PCT/EP 98/03204

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C04B38/00 C04B35/52

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 C04B B01J B01D H01G

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X, P	WO 97 20333 A (ALFAR INTERNATIONAL) 5 June 1997 see claims 1,3,6-8,14; table 1	1,2,5, 15-17, 19,20,24
A	---	3,4,6-8
X	DATABASE WPI Week 8945 Derwent Publications Ltd., London, GB; AN 89-325039 XP002079250 & DD 268 677 A (VEB LEUNA WERKE ULBRICHT) see abstract	19-22
X	FR 2 390 381 A (LE CARBONE-LORRAINE SA) 8 December 1978 see claims 1,2 -/--	19

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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Date of the actual completion of the international search

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Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Daeleman, P

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 98/03204

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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International Application No

PCT/EP 98/03204

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